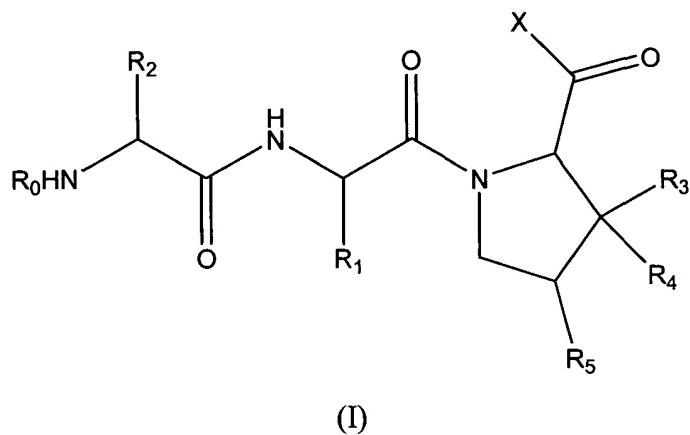


This listing of claims will replace all prior versions, and listings, of claims in the application:

**Listing of Claims**

1. (currently amended) Use of the compounds of the following formula (I): A method for the treatment of neurodegenerative diseases comprising administering an effective amount of a compound of formula (I):



wherein X represents OH, (C<sub>1-5</sub>) alkoxy, NH<sub>2</sub>, NH-C<sub>1-5</sub>-alkyl, or N(C<sub>1-5</sub> alkyl)<sub>2</sub>;

R<sub>1</sub> is a residue derived from one of the amino acids Phe, Tyr, Trp, Pro, which each may be optionally substituted with one or more (C<sub>1-5</sub>) alkoxy groups, (C<sub>1-5</sub>) alkyl groups or halogen atoms, as well as Ala, Val, Leu or Ile;

R<sub>2</sub> is a residue derived from one of the amino acids Gly, Ala, Ile, Val, Ser, Thr, Leu and or Pro;

Y<sub>1</sub> and Y<sub>2</sub> independently from each other represent H or (C<sub>1-5</sub>) alkyl;

R<sub>3</sub> and R<sub>4</sub> independently from each other represent H, OH, (C<sub>1-5</sub>) alkyl or (C<sub>1-5</sub>) alkoxy, provided that R<sub>3</sub> and R<sub>4</sub> are not both OH or (C<sub>1-5</sub>) alkoxy; and

R<sub>5</sub> represents H, OH, (C<sub>1-5</sub>) alkyl or (C<sub>1-5</sub>) alkoxy;  
or a pharmaceutically acceptable salt thereof;.  
~~for the preparation of a medicament useful in the treatment of neurodegenerative diseases.~~

2. (currently amended) The use method according to claim 1, wherein x represents (C<sub>1-5</sub>) alkoxy, NH<sub>2</sub>, NH-C<sub>1-5</sub>-alkyl, or N(C<sub>1-5</sub> alkyl)<sub>2</sub>.

3. (currently amended) The use method according to claim 1 or 2, wherein R<sub>3</sub> and R<sub>4</sub> independently from each other represent H, (C<sub>1-5</sub>) alkyl or (C<sub>1-5</sub>) alkoxy, provided that R<sub>3</sub> and R<sub>4</sub> are not (C<sub>1-5</sub>) alkoxy.

4. (currently amended) The use method according to ~~any of the previous claims~~ claim 1, wherein R<sub>5</sub> represents H, (C<sub>1-5</sub>) alkyl or (C<sub>1-5</sub>) alkoxy.

5. (currently amended) The use method according to ~~any of the previous claims~~ claim 1, wherein the neurodegenerative disease is Alzheimer's disease.

6. (currently amended) The use method according to ~~any of the previous claims~~ claim 1, wherein the neurodegenerative disease is mild cognitive impairment.

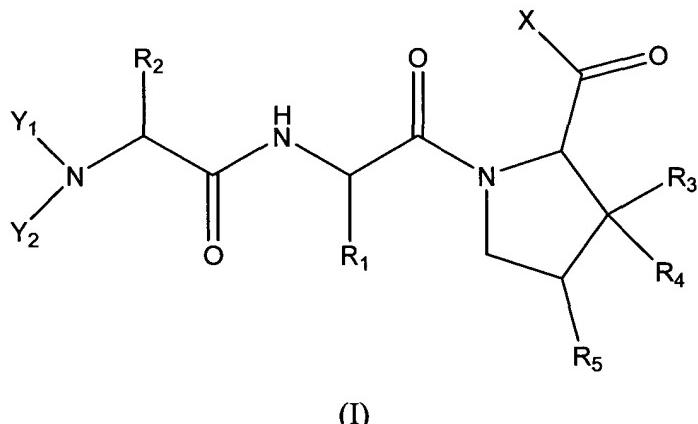
7. (currently amended) The use method according to ~~any of the previous claims~~ claim 1, wherein R<sub>1</sub> is a residue which is derived from one of the amino acids Phe, Tyr, Trp, each of which may optionally be substituted with a (C<sub>1-5</sub>) alkoxy group, a (C<sub>1-5</sub>) alkyl group or a halogen atom or which is derived from Ile.

8. (currently amended) The use method according to claim 7, wherein R<sub>1</sub> is a residue which is derived from Phe, which may optionally be substituted with a (C<sub>1-5</sub>) alkoxy group, a (C<sub>1-5</sub>) alkyl group or a halogen atom.

9. (currently amended) The use method according to any of the previous claims claim 1, wherein R<sub>2</sub> is a residue which is derived from the amino acid Gly or Ile.

10. (currently amended) The use method according to any of the previous claims claim 1, wherein the compound of formula (I) is glycyl-L-phenylalanyl-L-prolineamide, N,N-diethyl-isoleucyl-phenylalanyl-L-proline ethylamide, N,N-diethyl-isoleucyl-isoleucyl-prolineamide or a pharmaceutically acceptable salt thereof.

11. (currently amended) A pharmaceutical Pharmaceutical composition comprising compounds of the following formula (I):



wherein X represents OH, (C<sub>1-5</sub>) alkoxy, NH<sub>2</sub>, NH-C<sub>1-5</sub>-alkyl, N(C<sub>1-5</sub> alkyl)<sub>2</sub>;

R<sub>1</sub> is a residue derived from one of the amino acids Phe, which each may be optionally substituted with one or more (C<sub>1-5</sub>) alkoxy groups, (C<sub>1-5</sub>) alkyl groups or halogen atoms;

R<sub>2</sub> is a residue derived from one of the amino acids Gly, Ala, Ile, Val, Ser, Thr, Leu and Pro;

Y<sub>1</sub> and Y<sub>2</sub> independently from each other represent H or (C<sub>1-5</sub>) alkyl;

R<sub>3</sub> and R<sub>4</sub> independently from each other represent H, OH, (C<sub>1-5</sub>) alkyl or (C<sub>1-5</sub>)alkoxy, provided that R<sub>3</sub> and R<sub>4</sub> are not both OH or (C<sub>1-5</sub>) alkoxy; and

R<sub>5</sub> represents H, OH, (C<sub>1-5</sub>) alkyl or (C<sub>1-5</sub>) alkoxy;

or a pharmaceutically acceptable salt thereof ;

and pharmaceutically acceptable excipients.

12. (currently amended) The pharmaceutical Pharmaceutical composition according to claim 11, wherein x represents (C<sub>1-5</sub>) alkoxy, NH<sub>2</sub>, NH-C<sub>1-5</sub> alkyl or N(C<sub>1-5</sub> alkyl)<sub>2</sub>.

13. (currently amended) The pharmaceutical composition according to claims 11 or 12, wherein R<sub>2</sub> is a residue which is derived from the amino acid Gly.

14. (currently amended) The pharmaceutical Pharmaceutical composition according to ~~claims claim 11 to 13~~, wherein the compound of formula (I) is glycyl-L-phenylalanyl-L-prolineamide, N,N-diethyl-isoleucylphenylalanyl-L-proline ethylamide, N,N-diethyl-isoleucylisoleucyl-prolineamide or a pharmaceutically acceptable salt thereof.

15. (canceled)